

## DESCRIPTION

**Summary** TruCytes™ Potency CD19 Kit synthetic cells are lyophilized cell mimics that feature selected biomarkers for use in CAR-T potency assays. This product contains no bio-hazardous material, so it is safe to use in any environment and requires no special disposal.  
**For Research Use Only. Not for use in diagnostic or therapeutic procedures.**

## PRODUCT DETAILS

**Materials** TruCytes™ Potency CD19 are hydrogels that are in lyophilized form

**Handling and Safety** This product contains no bio-hazardous material, so it is safe to use in any environment and requires no special disposal.

**Shipping condition** The product is shipped on ice.

**Storage** Store lyophilized products at -20°C upon receipt.  
The product is recommended to be used immediately upon reconstitution.

**Expiration** 1 year from the date of manufacturing in lyophilized form

**Concentration** 2M particles/vial

**Product Format** 3 sealed glass vials per kit

**QC** IFN $\gamma$  release from CAR-T

## PREPARATION

**Reconstitution** Reconstitute in desired buffer or media.

**General Instructions for Use**

1. Tap down the vial to ensure that all cell mimics are collected at the bottom of the vial.
2. Add 1000  $\mu$ L of media to the vial and transfer contents to 15mL conical tube. Add additional 1000  $\mu$ L of media to vial, incubate 30 seconds to one minute, and transfer to same 15mL conical tube. Pellet cell mimics at 3000xg 3 mins room temperature.
3. Carefully remove supernatant from 1.5ml tube with 1000  $\mu$ L pipette. Resuspend pellet in 1000  $\mu$ L media, pellet as in (2).
4. Repeat step 3.
5. Carefully remove media and resuspend pellet in 1000  $\mu$ L media. Remove sample and count particles using a flow cytometer, hemacytometer, or automated counting instrument that incorporates phase contrast imaging to identify cells.
  - a. Note: If counts are performed on automated cell counter instrument with trypan blue exclusion, use total cell count. Automated cell counter instruments that rely exclusively on dye-based detection will require alternate methodology. Please reach out to Slingshot Biosciences for instrument-specific protocols or to discuss alternative options.
6. Pellet remaining sample as in (2), carefully remove supernatant, and resuspend mimics at desired concentration for assay.

DATA

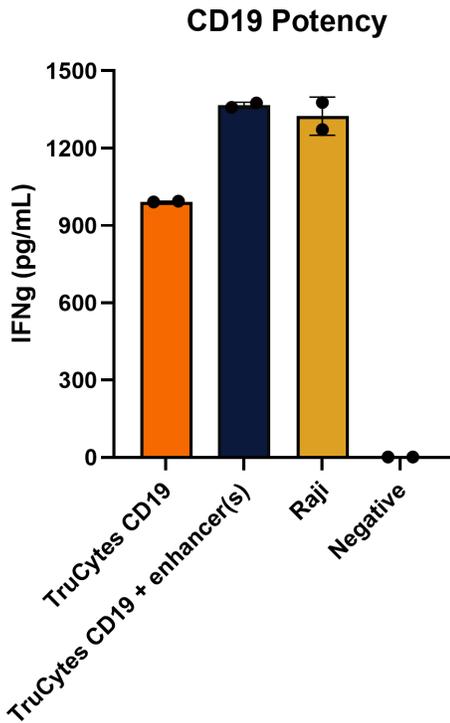


Figure 1. **Specific activation of CD19 CAR-T.** CD19 CAR-T were cultured at 1:1 E:T with TruCytes Potency negative (SL-00039), CD19 (SL-00037), and CD19 + enhancer(s) (SL-00038), or Raji cell line. Supernatant collected after 18 hours of culture was quantified for IFNγ with BD CBA on Cytex Aurora.